EXHIBIT 1

Orange Book: Approved Drug Products with Therapeutic Equivalence Evaluations

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U=HTTPS://WWW.ACCESSDATA.FDA.GOV/SCRIPTS/CDER/OB/RESULTS_PRODUCT.CFM?APPL_TYPE=A&APPL_NO=062337)

🐦 TWEET (HTTPS://TWITTER.COM/INTENT/TWEET/?TEXT=ORANGE BOOK: APPROVED DRUG PRODUCTS WITH THERAPEUTIC
EQUIVALENCE EVALUATIONS&URL=HTTPS://WWW.ACCESSDATA.FDA.GOV/SCRIPTS/CDER/OB/RESULTS_PRODUCT.CFM?
APPL_TYPE=A&APPL_NO=062337)

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EVALUATIONS&BODY=HTTPS://WWW.ACCESSDATA.FDA.GOV/SCRIPTS/CDER/OB/RESULTS_PRODUCT.CFM?
APPL_TYPE=A&APPL_NO=062337)

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Product Details for ANDA 062337

Expand all

DOXYCYCLINE HYCLATE (DOXYCYCLINE HYCLATE) EQ 50MG BASE	Marketing Status: Prescription

Active Ingredient: DOXYCYCLINE HYCLATE
Proprietary Name: DOXYCYCLINE HYCLATE
Dosage Form; Route of Administration: CAPSULE; ORAL
Strength: EQ 50MG BASE
Reference Listed Drug: No
Reference Standard: No
TE Code: AB
Application Number: A062337
Product Number: 001
Approval Date: Mar 29, 1982
Applicant Holder Full Name: MYLAN PHARMACEUTICALS INC
Marketing Status: Prescription
**Patent and Exclusivity Information (patent_info.cfm?
Product_No=001&Appl_No=062337&Appl_type=A)**

DOXYCYCLINE HYCLATE (DOXYCYCLINE HYCLATE)
EQ 100MG BASE

Marketing Status: Prescription

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EQUIVALENCE EVALUATIONS&URL=HTTPS://WWW.ACCESSDATA.FDA.GOV/SCRIPTS/CDER/OB/RESULTS_PRODUCT.CFM?
APPL_TYPE=A&APPL_NO=062031)

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EVALUATIONS&BODY=HTTPS://WWW.ACCESSDATA.FDA.GOV/SCRIPTS/CDER/OB/RESULTS_PRODUCT.CFM?
APPL_TYPE=A&APPL_NO=062031)

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Product Details for ANDA 062031

Expand all

DOXYCYCLINE HYCLATE (DOXYCYCLINE HYCLATE)
EQ 50MG BASE   Marketing Status: Prescription

Active Ingredient: DOXYCYCLINE HYCLATE
Proprietary Name: DOXYCYCLINE HYCLATE
Dosage Form; Route of Administration: CAPSULE; ORAL
Strength: EQ 50MG BASE
Reference Listed Drug: No
Reference Standard: No
TE Code: AB
Application Number: A062031
Product Number: 002
Approval Date: Oct 13, 1982
Applicant Holder Full Name: ACTAVIS LABORATORIES FL INC
Marketing Status: Prescription
**Patent and Exclusivity Information (patent_info.cfm?
Product_No=002&Appl_No=062031&Appl_type=A)**

DOXYCYCLINE HYCLATE (DOXYCYCLINE HYCLATE)
EQ 100MG BASE   Marketing Status: Prescription

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Product Details for ANDA 062500

Expand all

DOXYCYCLINE HYCLATE (DOXYCYCLINE HYCLATE)
EQ 50MG BASE   Marketing Status: Prescription

Active Ingredient: DOXYCYCLINE HYCLATE
Proprietary Name: DOXYCYCLINE HYCLATE
Dosage Form; Route of Administration: CAPSULE; ORAL
Strength: EQ 50MG BASE
Reference Listed Drug: No
Reference Standard: No
TE Code: AB
Application Number: A062500
Product Number: 001
Approval Date: Sep 11, 1984
Applicant Holder Full Name: CHARTWELL LIFE SCIENCE LLC
Marketing Status: Prescription
Patent and Exclusivity Information (patent_info.cfm? Product_No=001&Appl_No=062500&Appl_type=A)

DOXYCYCLINE HYCLATE (DOXYCYCLINE HYCLATE)
EQ 100MG BASE

Marketing Status: Prescription

Orange Book: Approved Drug Products with Therapeutic Equivalence Evaluations

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Product Details for ANDA 062396

Expand all

DOXYCYCLINE HYCLATE (DOXYCYCLINE HYCLATE)
EQ 50MG BASE   Marketing Status: Prescription

Active Ingredient: DOXYCYCLINE HYCLATE
Proprietary Name: DOXYCYCLINE HYCLATE
Dosage Form; Route of Administration: CAPSULE; ORAL
Strength: EQ 50MG BASE
Reference Listed Drug: No
Reference Standard: No
TE Code: AB
Application Number: A062396
Product Number: 002
Approval Date: Nov 7, 1984
Applicant Holder Full Name: HIKMA INTERNATIONAL PHARMACEUTICALS LLC
Marketing Status: Prescription
Patent and Exclusivity Information (patent_info.cfm? Product_No=002&Appl_No=062396&Appl_type=A)

DOXYCYCLINE HYCLATE (DOXYCYCLINE HYCLATE)
EQ 100MG BASE

Marketing Status: Prescription

Orange Book: Approved Drug Products with Therapeutic Equivalence Evaluations

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Product Details for ANDA 062676

Expand all

DOXYCYCLINE HYCLATE (DOXYCYCLINE HYCLATE)
EQ 50MG BASE   Marketing Status: Prescription

Active Ingredient: DOXYCYCLINE HYCLATE
Proprietary Name: DOXYCYCLINE HYCLATE
Dosage Form; Route of Administration: CAPSULE; ORAL
Strength: EQ 50MG BASE
Reference Listed Drug: No
Reference Standard: No
TE Code: AB
Application Number: A062676
Product Number: 002
Approval Date: Jul 10, 1986
Applicant Holder Full Name: SUN PHARMACEUTICAL INDUSTRIES INC
Marketing Status: Prescription
Patent and Exclusivity Information (patent_info.cfm? Product_No=002&Appl_No=062676&Appl_type=A)

DOXYCYCLINE HYCLATE (DOXYCYCLINE HYCLATE)
EQ 100MG BASE

Marketing Status: Prescription